

CLAIMS

1. A method for inducing vasodilation and/or increasing blood flow in a subject, comprising administering to the subject an effective amount of a pharmaceutically-acceptable salt of nitrite for a sufficient period of time to induce vasodilation and/or increase blood flow in the subject.
2. The method of claim 1, wherein the pharmaceutically-acceptable salt of nitrite reacts in the presence of hemoglobin in the subject to release nitric oxide.
- 10 3. The method of claim 1, wherein the effective amount of the pharmaceutically-acceptable salt of nitrite:
 - induces production in the subject of no more than about 25% methemoglobin;
 - induces production in the subject of no more than about 20% methemoglobin;
 - induces production in the subject of no more than about 10% methemoglobin;
 - 15 induces production in the subject of no more than about 8% methemoglobin; or
 - induces production in the subject of no more than about 5% methemoglobin.
4. The method of claim 1, wherein the effective amount of the pharmaceutically-acceptable salt of nitrite induces production in the subject of no more than about 3% methemoglobin.
- 20 5. The method of claim 1, comprising administering sodium nitrite by injection at about 36 μ moles per minute for at least five minutes into the forearm brachial artery of the subject.
6. The method of claim 1, wherein the effective amount of the pharmaceutically-acceptable salt of nitrite is administered to a circulating concentration in the subject of about 0.6 to 240 μ M.
- 25 7. The method of any one of claims 1-6, wherein the pharmaceutically-acceptable salt of nitrite comprises as the cation sodium, potassium, or arginine.
- 30 8. The method of claim 7, wherein the nitrite is administered as sodium nitrite.
9. The method of any of claims 1-8, wherein the administration of the nitrite is parenteral, oral, bucal, rectal, *ex vivo*, or intraocular.
- 35 10. The method of any of claims 1-8, wherein the administration of the nitrite is peritoneal, intravenous, intraarterial, subcutaneous, inhaled, intramuscular, or into a cardiopulmonary bypass circuit.

11. The method of any one of claims 1-10, wherein the subject is a mammal.
12. The method of claim 11, wherein the subject is a human.
5
13. The method of any one of claims 1-12, wherein the nitrite is administered in combination with at least one additional agent.
14. The method of claim 13, wherein the additional agent is one or more selected from
10 the list consisting of penicillin, hydroxyurea, butyrate, clotrimazole, arginine, or a phosphodiesterase inhibitor.
14. The method of claim 14, wherein the phosphodiesterase inhibitor is sildenafil.
- 15 15. The method of any one of claims 1-13, wherein the subject has elevated blood pressure, and the method is a method for treating at least one vascular complication associated with
the elevated blood pressure.
16. The method of any one of claims 1-13, wherein the subject has a hemolytic
20 condition, and the method is a method for treating at least one vascular complication associated with
the hemolytic condition.
17. The method of claim 15 or 16, wherein the at least one vascular complication is one
or more selected from the group consisting of pulmonary hypertension, systemic hypertension,
25 peripheral vascular disease, trauma, cardiac arrest, general surgery, organ transplantation, cutaneous
ulceration, acute renal failure, chronic renal failure, intravascular thrombosis, angina, an ischemia-
reperfusion event, an ischemic central nervous system event, and death.
18. The method of claim 17, wherein the hemolytic condition is one or more selected
30 from the group consisting of sickle cell anemia, thalassemia, hemoglobin C disease, hemoglobin SC
disease, sickle thalassemia, hereditary spherocytosis, hereditary elliptocytosis, hereditary ovalcytosis,
glucose-6-phosphate deficiency and other red blood cell enzyme deficiencies, paroxysmal nocturnal
hemoglobinuria (PNH), paroxysmal cold hemoglobinuria (PCH), thrombotic thrombocytopenic
35 purpura/hemolytic uremic syndrome (TTP/HUS), idiopathic autoimmune hemolytic anemia, drug-
induced immune hemolytic anemia, secondary immune hemolytic anemia, non-immune hemolytic
anemia caused by chemical or physical agents, malaria, falciparum malaria, bartonellosis, babesiosis,
clostridial infection, severe haemophilus influenzae type b infection, extensive burns, transfusion
reaction, rhabdomyolysis (myoglobinemia), transfusion of aged blood, transfusion of hemoglobin,

transfusion of red blood cells, cardiopulmonary bypass, coronary disease, cardiac ischemia syndrome, angina, iatrogenic hemolysis, angioplasty, myocardial ischemia, tissue ischemia, hemolysis caused by intravascular devices, and hemodialysis.

5 19. The method of any one of claims 1-13, wherein the subject has a condition associated with decreased blood flow to a tissue, and the method is a method to increase blood flow to the tissue of the subject.

10 20. The method of claim 19, wherein the decreased blood flow to the tissue is caused directly or indirectly by at least one condition selected from the group consisting of: sickle cell anemia, thalassemia, hemoglobin C disease, hemoglobin SC disease, sickle thalassemia, hereditary spherocytosis, hereditary elliptocytosis, hereditary ovalcytosis, glucose-6-phosphate deficiency and other red blood cell enzyme deficiencies, paroxysmal nocturnal hemoglobinuria (PNH), paroxysmal cold hemoglobinuria (PCH), thrombotic thrombocytopenic purpura/hemolytic uremic syndrome (TTP/HUS), idiopathic autoimmune hemolytic anemia, drug-induced immune hemolytic anemia, secondary immune hemolytic anemia, non-immune hemolytic anemia caused by chemical or physical agents, malaria, falciparum malaria, bartonellosis, babesiosis, clostridial infection, severe haemophilus influenzae type b infection, extensive burns, transfusion reaction, rhabdomyolysis (myoglobinemia), transfusion of aged blood, transfusion of hemoglobin, transfusion of red blood cells, cardiopulmonary bypass, coronary disease, cardiac ischemia syndrome, angina, iatrogenic hemolysis, angioplasty, myocardial ischemia, tissue ischemia, hemolysis caused by intravascular devices, hemodialysis, pulmonary hypertension, systemic hypertension, cutaneous ulceration, acute renal failure, chronic renal failure, intravascular thrombosis, and an ischemic central nervous system event.

25 21. The method of claim 20, wherein the tissue is an ischemic tissue.

30 22. The method of any one of claims 19-21, wherein the tissue is one or more tissues selected from the group consisting of neuronal tissue, bowel tissue, intestinal tissue, limb tissue, lung tissue, central nervous tissue, or cardiac tissue.

23. The method of claim 15, wherein the elevated blood pressure comprises elevated blood pressure in the lungs.

35 24. The method of claim 23, wherein the subject has neonatal pulmonary hypertension.

25. The method of claim 23, wherein the subject has primary and/or secondary pulmonary hypertension.

26. The method of any of any one of claims 23-26, wherein the pharmaceutically-acceptable salt of nitrite is nebulized.

5 27. The method of claim 26, wherein the pharmaceutically-acceptable salt of nitrite is administered to a circulating concentration in the subject of:

no more than about 100 μ M;

no more than about 50 μ M;

no more than about 20 μ M;

10 no more than about 16 μ M; or

less than about 16 μ M.